# PREMIER VALUE NIGHTTIME SLEEP-AID LIQUID SLEEP-AID- diphenhydramine hydrochloride liquid Chain Drug Consortium

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**Premier Value Nighttime Sleep-Aid** 

**Drug Facts** 

# Active ingredients (in each 30 mL dose cup or 2 tablespoons)

Diphenhydramine HCl 50 mg

#### **Purpose**

Nighttime sleep-aid

#### Uses

- for the relief of occasional sleeplessness
- reduces time to fall asleep if you have difficulty falling asleep

## Warnings

#### Do not use

- for children under 12 years of age
- with any other product containing diphenhydramine, even one used on skin
- with other drugs that cause drowsiness such as antihistamines and nighttime cold/flu products

# Ask a doctor before use if you have

- · a breathing problem such as asthma, emphysema, or chronic bronchitis
- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- heart disease

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers or any other sleep aid

# When using this product

- avoid alcoholic beverages and other drugs that cause drowsiness
- · drowsiness will occur
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if sleeplessness persists continuously for more than 2

weeks. Insomnia may be a symptom of serious underlying medical illness.

**If pregnant or breast-feeding**, ask a health professional before use.

Keep out of reach of children.

#### Overdose warning

Taking more than directed can cause serious health problems. In case of overdose, get medical help or contact a Poison Control Center right away.

#### **Directions**

- take only one dose per day (24 hours) see Overdose warning
- use dose cup or tablespoon

adults & children 12 yrs & over	One Dose = 30 mL (2 tablespoons) at bed time if
	needed or as directed by a doctor

#### Other information

- each 30 mL dose (2 tablespoons) contains: sodium 23 mg
- store at room temperature
- protect from light. Does not meet USP <671>.

## Inactive ingredients

anhydrous citric acid, carboxymethylcelluose sodium, FD&C blue 1, FD&C red 40, flavor, glycerin, potassium citrate, purified water, sodium benzoate, sorbitol, sucralose.

# Failure to follow these warnings could result in serious consequences

**TAMPER EVIDENT:** This package is safety sealed & child resistant. Use only if blisters are intact. If difficult to open, use scissors.

This product is not manufactured or distributed by Procter & Gamble, the distributor of  $ZzzQuil^{m}$ .

Distributed by:

Chain Drug Consortium LLC.

3301 N.W. Boca Raton Blvd.

Suite 101, Boca Raton FL 33431

#### PRINCIPAL DISPLAY PANEL - 354 ml Bottle Label

Premier Value Nighttime

Sleep-Aid

### Diphenhydramine HCI /

Non-Habit Forming

Berry Flavor

Not for treating Cold or Flu See Warnings

12 FL OZ (354 ml)



# PREMIER VALUE NIGHTTIME SLEEP-AID LIQUID SLEEP-AID

diphenhydramine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-219
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
<b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	50 mg in 30 mL		

Inactive Ingredients		
Ingredient Name	Strength	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		

GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	PURPLE	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

l	Packaging				
	# Item	Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:680 219-12		354 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/10/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M010	12/10/2012	

# Labeler - Chain Drug Consortium (101668460)

Revised: 11/2023 Chain Drug Consortium